

# Origin of the Helminth Community of an Exotic Invasive Lizard, the Brown Anole, *Anolis sagrei* (Squamata: Polychrotidae), in Southwestern Taiwan<sup>1</sup>

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**Abstract:** Composition of the helminth community of the brown anole, *Anolis sagrei*, an exotic invasive species in Taiwan, was studied to identify the emigration point of this lizard. A total of 5,757 helminths was found, of which 5,734 (99.6%) were the nematode *Cyrtosomum penneri*. Also found were the digenean *Mesocoelium monas* (21, 0.4%) and one each of the nematodes *Parapharyngodon* sp. (female) and Acuariidae gen. sp. (larva). *Cyrtosomum penneri* has previously been reported in *A. sagrei* in Florida, supporting the contention that the Taiwan population of *A. sagrei* originated from Florida. This report provides a basis upon which future *A. sagrei* parasite studies in Taiwan can be based, and a helminth list for *A. sagrei* is included for future reference.

THE BROWN ANOLE, *Anolis sagrei* Duméril and Bibron, 1837, is native to Cuba, the Bahamas, and adjacent islands (Henderson and Powell 2009). It is a widespread and successful colonizer, and Kraus (2009) published a list of invaded sites, suggesting Hawai'i and Taiwan as more recent sites of colonization. The Hawaiian (O'ahu) population was first noticed in 1980 (McKeown 1996) and is most likely of Florida or Caribbean origin (Goldberg and Bursey 2000). The Taiwan population was first noted in 2000 and because the highest concentration occurred in the immediate vicinity of a plant nursery (Norval et al. 2002),

it is likely that these lizards were introduced with nursery products. The effects of parasite introductions by invasive reptiles merit investigation as part of the ecological and socioeconomic impact of naturalized animals (Lever 2003). Introduced brown anoles are considered to have originated from Florida (Kolbe et al. 2004) and have transported helminth parasites of Florida/Caribbean distributions to Hawai'i (Goldberg and Bursey 2000). Identification of the helminths present in introduced reptiles is important in tracing emigration pathways. The purpose of the study reported here was to examine the helminth population harbored by *A. sagrei* in Taiwan to identify their emigration site and to provide a basis upon which future *A. sagrei* parasite studies in Taiwan can be based.

## MATERIALS AND METHODS

Between 20 September 2007 and 19 September 2008, 347 brown anoles (175 male, snout-vent length [SVL] = 24–60 mm; 172 female, SVL = 22–47 mm) were collected monthly by hand at night in agricultural fields, a cemetery, and along roads in Santzepu, Sheishan District, Chiayi County, Taiwan (23° 25' 51" N, 120° 28' 30" E). Due to typhoon Kalmaegi, no sample was obtained for July 2008. At the point of capture each specimen was allocated a field number. The following morning, each lizard was measured to the nearest millimeter

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TABLE 1

Number, Prevalences (%), Mean Intensity  $\pm 1$  SD, Range, and Abundance for Helminths of *Anolis sagrei* from Taiwan

Helminth	<i>n</i>	Prevalence	Mean Intensity	Range	Abundance
Digenea					
<i>Mesocoelium monas</i>	21	3.7	1.6 $\pm$ 1.2	1–5	0.061
Nematoda					
<i>Cyrtosomum penneri</i>	5,734	37.7	43.7 $\pm$ 50.3	1–265	16.524
<i>Parapharyngodon</i> sp.	1	0.3	1	—	0.003
Acuariid larva	1	0.3	1	—	0.003

with a transparent plastic ruler, then killed with ether, and fixed by injecting 10% formalin into the body cavity. After a 2-day fixation period, each specimen was placed individually in a sealed plastic bag filled with 75% alcohol. All specimens were shipped to Whittier College, Whittier, California, where the body cavity of each lizard was opened by a longitudinal incision from throat to vent and the gastrointestinal tract was removed by cutting across the esophagus and rectum. The esophagus, stomach, and small and large intestines were slit longitudinally and separately searched for helminths under a dissecting microscope. The wall of the digestive tract was also examined for cysts containing larval helminths. For study, helminths were cleared in a drop of undiluted glycerol on a glass slide. Nematodes were identified from these temporary preparations; digeneans were regressively stained in hematoxylin, cleared in xylene, and mounted in Canada balsam. Identifications are based upon the reference keys of Anderson et al. (1974) and Prudhoe and Bray (1982). Population terminology is according to Bush et al. (1997). Statistical analysis followed the methods outlined by Brower et al. (1998). The anoles were deposited in the herpetology collection of the Natural History Museum of Los Angeles County (LACM), Los Angeles, California. Selected helminths were deposited in the United States Parasite Collection (USNPC), Beltsville, Maryland, as *Mesocoelium monas* (USNPC 101987, 101988), *Cyrtosomum penneri* (USNPC 101989), *Parapharyngodon* sp. (USNPC 101990), and Acuariidae gen. sp. (USNPC 101991).

## RESULTS

One hundred and thirty seven (39%) anoles harbored a total of 5,757 helminths. Four species of helminths were found: one digenean species, *Mesocoelium monas* (Rudolphi, 1819) Freitas, 1957 (small intestine); and three nematode species, *Cyrtosomum penneri* Gambino, 1957 (large and small intestines), an unidentified species of *Parapharyngodon* (large intestine), and a larva of an acuariid nematode (small intestine). The number, prevalence (number of infected host species as percentage), mean intensity (mean number of helminths per infected host), and abundance (total number of parasites in a host sample divided by total number of hosts in the sample) by helminth species are given in Table 1. Other reported helminths from *A. sagrei* are listed in Table 2. Of the 5,757 helminths found in *A. sagrei* from Taiwan, 5,734 (99.6%) were individuals assigned to *C. penneri* based on the spicule appearance. There was no significant difference in infection rate of male and female anoles by *C. penneri* ( $\chi^2 = 3.18$ , ddf = 1,  $P > .05$ ); thus, no subsequent separation of data by host gender was made. The numbers of individuals of the other identified helminth species are too few for analysis by gender. For *C. penneri*, prevalence by month is given in Figure 1, and abundance (total number of parasite individuals/total number of hosts) by month is given in Figure 2. One hundred and thirty one (96% of infected hosts) anoles were infected by *C. penneri*; 13 anoles (9%) by *M. monas*, one anole (<1%) by a *Parapharyngodon* sp., and one anole by an

TABLE 2  
Reported Helminths of *Anolis sagrei*

Helminth	Locality	Prevalence	Reference
<b>Acanthocephala</b>			
<i>Acanthocephalus bufonis</i>	Hawai'i	2/62	Goldberg and Bursey (2000)
	Hawai'i	8/281	Goldberg et al. (2002)
<i>Centrorhynchus</i> sp.	Bahamas	1/12	Goldberg et al. (1994)
<b>Eucestoda</b>			
<i>Oochoristica</i> sp.	Hawai'i	8/281	Goldberg et al. (2002)
<b>Nematoda</b>			
<i>Abbreviata</i> sp. <sup>a</sup>	Cuba	12/138	Coy Otero and Barus (1979)
Acuariidae <sup>a</sup>	Hawai'i	66/281	Goldberg et al. (2002)
<i>Atractis opeatura</i>	Cuba	2/138	Coy Otero and Barus (1979)
<i>Atractis scolopori</i>	Hawai'i	39/62	Goldberg and Bursey (2000)
	Hawai'i	189/281	Goldberg et al. (2002)
<i>Cyrtosomum penneri</i>	Florida	22/25	Goldberg et al. (1994)
<i>Cyrtosomum scolopori</i>	Bahamas	23/45	Goldberg et al. (1994)
	Cuba	4/21	Barus and Coy Otero (1969)
	Cuba	4/21	Coy Otero (1970)
	Cuba	4/19	Coy Otero and Barus (1973)
	Cuba	33/138	Coy Otero and Barus (1979)
	Florida	36/100	Price and Underwood (1984)
<i>Oswaldocruzia lenteixeirai</i>	Bahamas	3/45	Goldberg et al. (1994)
	Cuba	3/138	Coy Otero and Barus (1979)
<i>Ozolaimus monbystera</i>	Cuba	2/138	Coy Otero and Barus (1979)
<i>Parapharyngodon cubensis</i>	Bahamas	1/45	Goldberg et al. (1994)
	Cuba	1/21	Barus and Coy Otero (1969)
	Cuba	1/21	Coy Otero (1970)
	Cuba	2/138	Coy Otero and Barus (1979)
<i>Physaloptera squamatae</i>	Bahamas	20/45	Goldberg et al. (1994)
	Cuba	3/138	Coy Otero and Barus (1979)
	Florida	50/100	Price and Underwood (1984)
	Florida	15/25	Goldberg et al. (1994)
	Hawai'i	61/62	Goldberg and Bursey (2000)
	Hawai'i	165/281	Goldberg et al. (2002)
Physalopteridae gen. sp.	Cuba	1/138	Coy Otero and Barus (1979)
<i>Physocephalus</i> sp. <sup>a</sup>	Hawai'i	50/62	Goldberg and Bursey (2000)
	Hawai'i	145/281	Goldberg et al. (2002)
<i>Porrocaecum</i> sp. <sup>a</sup>	Bahamas	1/45	Goldberg et al. (1994)
	Cuba	5/138	Coy Otero and Barus (1979)
<i>Skrjabinoptera pbrynosoma</i>	Cuba	7/21	Barus and Coy Otero (1969)
	Cuba	7/21	Coy Otero (1970)
	Cuba	55/138	Coy Otero and Barus (1979)
<i>Strongyloides</i> sp.	Bahamas	1/45	Goldberg et al. (1994)
<i>Trichospirura teixeirai</i>	Cuba	13/138	Coy Otero and Barus (1979)
<b>Pentastomida</b>			
<i>Kiricephalus pattoni</i> <sup>b</sup>	Taiwan	Not stated	Norval et al. (2009)
<i>Raillietiella frenatus</i>	Hawai'i	8/62	Goldberg and Bursey (2000)
	Hawai'i	23/281	Goldberg et al. (2002)
<b>Trematoda</b>			
<i>Mesocoelium monas</i>	Florida	10/100	Price and Underwood (1984)
	Florida	2/82	Sellers and Graham (1987)
	Hawai'i	3/62	Goldberg and Bursey (2000)
<i>Platynosomum fastosum</i> <sup>a</sup>	Florida	Not stated	Eckerlin and Leigh (1962)
	Hawai'i	2/62	Goldberg and Bursey (2000)
<i>Urotrema scabridum</i>	Bahamas	3/45	Goldberg et al. (1994)
	Cuba	5/21	Coy Otero (1970)
	Florida	7/82	Sellers and Graham (1987)
	Florida	4/25	Goldberg et al. (1994)
<i>Urotrema wardi</i>	Cuba	2/21	Coy Otero (1970)

<sup>a</sup> Larvae.

<sup>b</sup> Nymph.

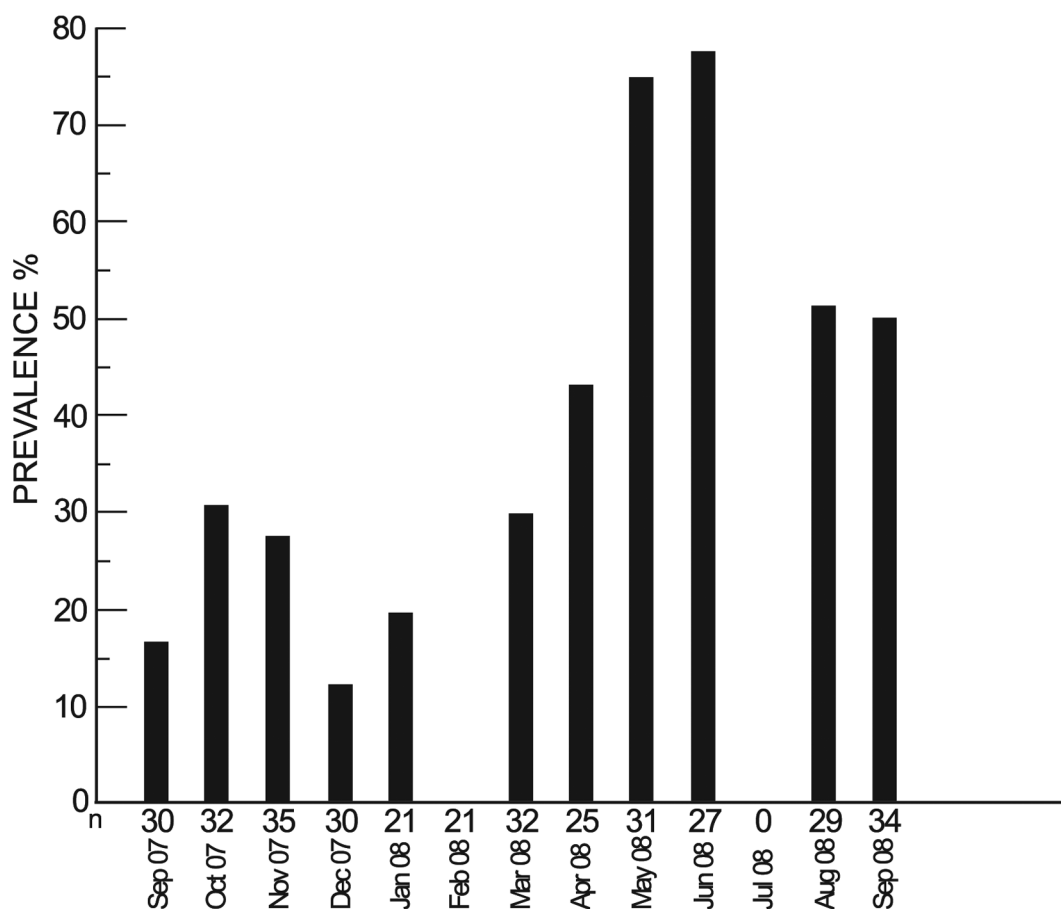


FIGURE 1. Prevalence by month of *Cyrtosomum penneri*. Number of hosts examined appears below axis.

acuariid larva; eight anoles had concurrent infections of *C. penneri* and *M. monas*; one anole had a concurrent infection of *C. penneri* and *Parapharyngodon* sp. Mean number of helminths harbored per infected host was  $42 \pm 50$  (1–265).

#### DISCUSSION

As noted in Table 2, two species of *Cyrtosomum* have been reported in *A. sagrei* from Florida: *C. penneri* by Goldberg et al. (1994) and *C. scelopori* Gedoelst, 1919, by Price and Underwood (1984). *Cyrtosomum penneri* can be distinguished from *C. scelopori* by the spicule appearance: the spicules of *C. scelopori* are

equal in length (Gedoelst 1919), and those of *C. penneri* are unequal in length (Gambino 1957). We have assigned our specimens to *C. penneri* because the males possess spicules of unequal length. The life cycle of *C. penneri* has not been studied; however, eggs of atractids hatch and larvae develop to third stage in utero; thus, autoinfection is the rule (Anderson 2000). Petter (1966) postulated that in tortoises, at least, atractids are transmitted during copulation. It is of interest to note that in collared lizards, *Crotaphytus collaris* (Say, 1823), infection rates of *C. penneri* were significantly higher in sexually mature individuals than in juveniles (Pfaffenberger et al. 1986), and in our study the smallest infected male

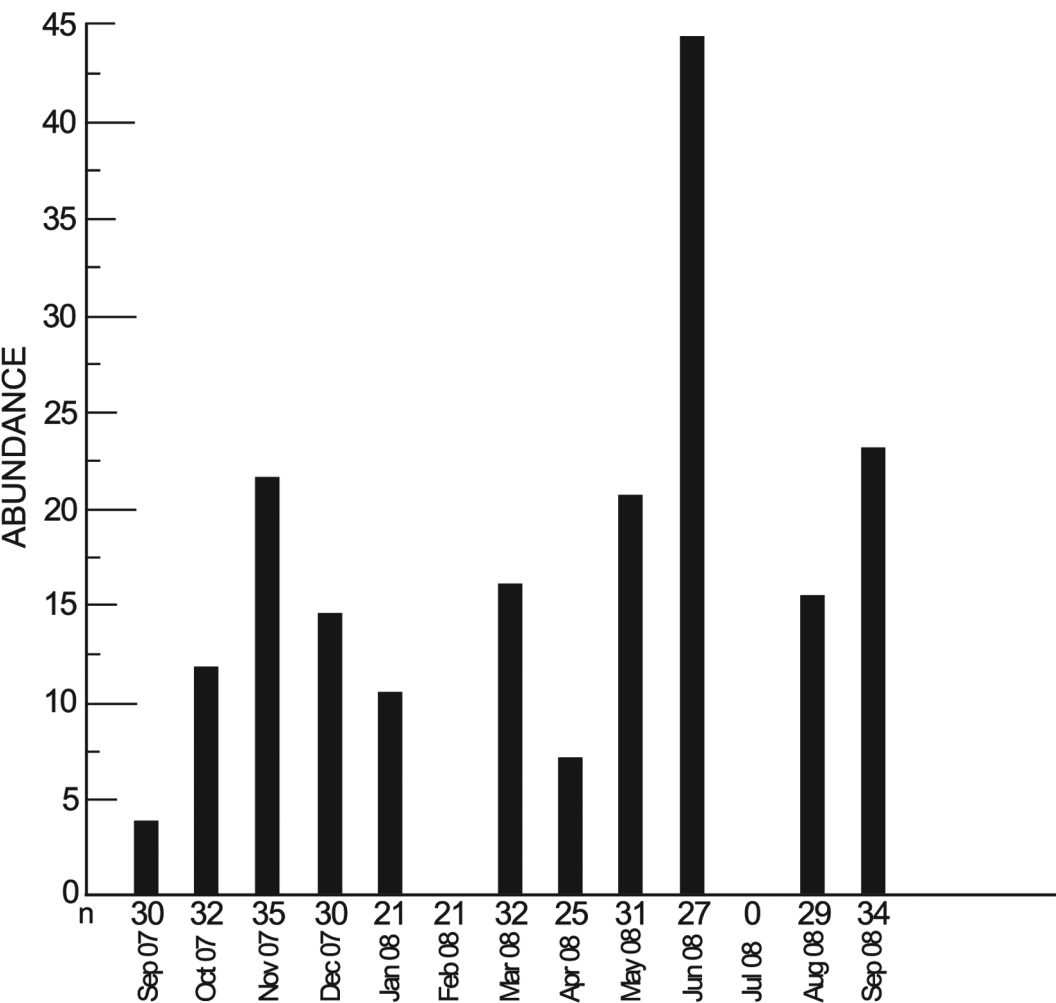


FIGURE 2. Abundance by month of *Cyrtosomum penneri*. Number of hosts examined appears below axis.

was 34 mm SVL (infected individuals, mean  $50 \pm 6$  mm, range 34–60), and the smallest infected female was 35 mm SVL (infected individuals, mean  $41 \pm 3$  mm, range 35–48); all, except the smallest infected male, were within the breeding size range (see Goldberg et al. 2002) and thus supportive of copulation as a possible method for new infections of *C. penneri*. Henderson and Powell (2009) reported a life expectancy of 1.0 yr for *A. sagrei* males and 1.8 yr for females. Thus, maturing uninfected anoles, and seasonal reproduction, could combine to produce conditions that

favor low prevalence in winter and high prevalence in summer so that by the end of summer almost all breeding anoles are infected (Figure 1). A similar pattern is seen with abundance (Figure 2). There is lower abundance in winter, and greater abundance in summer; however, abundance of *C. penneri* is indicative of the fecundity of their infrapopulations within *A. sagrei* hosts and the time elapsed since copulation by hosts. Conceivably, the greatest intensity for *C. penneri* could be just before death of the *A. sagrei* host, but without consequence to the host population.

Baer (1951) suggested that these nematodes, which occur in large numbers and in all stages of development, are possibly living on partially digested matter and should be considered as commensals rather than true parasites.

*Mesocoelium monas* is a parasite of the intestines of amphibians and reptiles in tropical and subtropical regions (Prudhoe and Bray 1982). Fischthal and Kuntz (1975) reported *M. monas* (= *Mesocoelium sociale* Lühe, 1901) from two species of amphibians, *Fejervarya limnocharis* (Gravenhorst, 1829) and *Duttaphrynus melanostictus* (Schneider, 1799); a lizard species, *Japalura swinhonis* Günther, 1864; and a species of snake, *Amphiesma stolatum* (Linnaeus, 1758), all from Taiwan and very common in the locality where our *A. sagrei* were collected. Price and Underwood (1984) and Sellers and Graham (1987) reported *M. monas* from *A. sagrei* collected in Florida. But it should also be noted that *M. monas* occurs in all biogeographic realms except the Palearctic (Burse and Goldberg 2003). Thus, no assumption as to whether *M. monas* was transported or acquired can be made. Infection by *M. monas* occurs through ingestion of infected molluscan intermediate hosts or vegetation contaminated with cercarial cysts (Prudhoe and Bray 1982).

There was one female individual of a *Parapharyngodon* sp. in our sample. To our knowledge, there are no reports of a species of *Parapharyngodon* in any host from Taiwan, nor are there reports of *Parapharyngodon* in *A. sagrei* from Florida. However, there are reports of *Parapharyngodon cubensis* (Barus and Coy Otero, 1969) in *A. sagrei* from the Bahamas and Cuba (Barus and Coy Otero 1969, Coy Otero 1970, Coy Otero and Barus 1979, Goldberg et al. 1994). It should be noted that Hasegawa (1988) reported an unidentified species of *Parapharyngodon* in a lizard, *Ateuchosaurus pellopleurus* (Hallowell, 1861), collected on Okinawa. Species of *Parapharyngodon* are identified utilizing characteristics of male individuals; thus identification of our specimen to the species level is not possible.

One larva of a species of acuariid nematode was found. Members of the Acuariidae are usually parasites of the alimentary tract of birds and use a variety of invertebrates as in-

termediate hosts, so lizards are considered paratenic/accidental hosts (Anderson 2000).

It is becoming evident that immigrant lizards, under favorable conditions, may transport helminths. Ecological conditions favoring egg survival for monoxenous helminths (direct life cycle, no intermediate host utilized) and appropriate intermediate hosts for heteroxenous helminths (indirect life cycle, obligatory intermediate host) are important considerations in the establishment of immigrant helminth populations. It is evident that the population of *C. penneri* is well established and persistent. But the question becomes, especially for autoinfective helminths, what conditions must be met to spread throughout the host community? More work remains to determine if *C. penneri* can become established in native Taiwanese lizards, as well as if *A. sagrei* will acquire any of the native parasites.

A study by Kolbe et al. (2004) showed that introduced *A. sagrei* from Hawai'i and Taiwan are considered to have originated from Florida but could not indicate if the introductions were related. The dominant nematodes from the studies in Hawai'i, *Cyrtosomum* (as *Atractis*) *scelopori* and *Physaloptera squamatae* Harwood, 1932 (Goldberg and Bursey 2000), are absent from lizards in the study reported here, and the dominant nematode from this study, *C. penneri*, was not recorded in Hawai'i, so it seems that the introduction of *A. sagrei* in these two localities were isolated incidents. That the Taiwan population of *A. sagrei* originated in Florida is given additional support by the fact that all three helminths, *C. penneri*, *C. scelopori*, and *P. squamatae*, occur in Florida (Table 2).

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